



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁷ :C12N 15/12, C07K 14/705, 14/72, A61K
38/04, A61P 15/06, G01N 33/50, 33/68

A1

(11) International Publication Number:

WO 00/17348

(43) International Publication Date:

30 March 2000 (30.03.00)

(21) International Application Number: PCT/CA99/00844

(22) International Filing Date: 15 September 1999 (15.09.99)

(30) Priority Data:

09/154,627

17 September 1998 (17.09.98) US

(63) Related by Continuation (CON) or Continuation-in-Part (CIP) to Earlier Application

US

09/154,627 (CIP)

Filed on

17 September 1998 (17.09.98)

(71) Applicant (for all designated States except US): HOPITAL
SAINTE-JUSTINE [CA/CA]; 3175 Côte Sainte-Catherine,
Montréal, Québec H3T 1C5 (CA).

(72) Inventors; and

(75) Inventors/Applicants (for US only): CHEMTOB, Sylvain
[CA/CA]; 6885 Banting, Montréal, Québec H4W 1G1 (CA).
PERI, Krishna, G. [CA/CA]; #315, 3555 Atwater Avenue,
Montréal, Québec H3H 1Y3 (CA).(74) Agents: CÔTÉ, France et al.; Swabey Ogilvy Renault, Suite
1600, 1981 McGill College Avenue, Montréal, Québec H3A
2Y3 (CA).(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG,
BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE,
ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP,
KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD,
MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD,
SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US,
UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS,
MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ,
BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE,
CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC,
NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA,
GN, GW, ML, MR, NE, SN, TD, TG).

Published

With international search report.

Before the expiration of the time limit for amending the
claims and to be republished in the event of the receipt of
amendments.

(54) Title: G PROTEIN-COUPLED RECEPTOR AGONISTS OR ANTAGONISTS

(57) Abstract

The present invention relates to a new class of G protein-coupled receptor agonist or antagonist, which specifically binds to the receptor protein structural elements, thus altering signal transmission and subsequent physiological effects. Described herein are peptide sequences derived from the G protein-coupled receptor protein, produced by chemical methods as selective inhibitors of signal transduction associated with stimulation of the receptor by its ligand. Such peptides or molecules derived from their primary, secondary or tertiary structures may be used as effective tocolytics for the prevention of premature labor or be used for the treatment of dysmenorrhea.

Fold stimulation of
IP hydrolysis